

**PCT**WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>6</sup> :</b> <b>A61K 7/48, 7/035</b>	<b>A1</b>	<b>(11) International Publication Number:</b> <b>WO 98/00104</b> <b>(43) International Publication Date:</b> 8 January 1998 (08.01.98)
<b>(21) International Application Number:</b> PCT/EP97/03063 <b>(22) International Filing Date:</b> 10 June 1997 (10.06.97) <b>(30) Priority Data:</b> 60/020,748 28 June 1996 (28.06.96) US <b>(71) Applicant (for AU BB CA GB GH IE IL KE LC LK LS MN MW NZ SD SG SZ TT UG only):</b> UNILEVER PLC [GB/GB]; Unilever House, Blackfriars, London EC4P 4BQ (GB). <b>(71) Applicant (for all designated States except AU BB CA GB GH IE IL KE LC LK LS MN MW NZ SD SG SZ TT UG):</b> UNILEVER N.V. [NL/NL]; Weena 455, NL-3013 AL Rotterdam (NL). <b>(72) Inventors:</b> DOBKOWSKI, Brian, John; 31 Wakeley Street, Shelton, CT 06484 (US). ZNAIDEN, Alexander, Paul; 110 Fox Road, Trumbull, CT 06611 (US). CHENEY, Michael, Charles; 383 Sigwin Drive, Fairfield, CT 06430 (US). <b>(74) Agent:</b> ROTS, Maria, Johanna, Francisca; Unilever PLC, Patent Division, Colworth House, Sharnbrook, Bedford MK44 1LQ (GB).		<b>(81) Designated States:</b> AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title:</b> POWDERED COSMETIC COMPOSITIONS CONTAINING SILICONE ELASTOMERS <b>(57) Abstract</b>  A cosmetic powder is provided which includes a cross linked non-emulsifying siloxane elastomer, a powdered inorganic material and a skin treatment agent. Inclusion of the elastomer allows for the coupling of water as well as water soluble salts into the cosmetic powder.		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CJ	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

- 1 -

POWDERED COSMETIC COMPOSITIONS  
CONTAINING SILICONE ELASTOMERS

5

BACKGROUND OF THE INVENTION

10

Field of the Invention

The invention concerns cosmetic compositions in powdered form for topical application to the skin having improved aesthetics through use of special silicone elastomers.

15

The Related Art

20

Consumers are ever more demanding in the aesthetics delivered by such powdered cosmetics as talcum powder. Much has already been done to improve these products. For instance, Vaseline® Intensive Care® Baby Powder besides talc includes fragrance and the skin conditioning agents of petrolatum and methicone.

25

30

The present invention has sought a system which allows inorganic powders to store a high level of hydrophobic and/or hydrophilic liquids. It is of interest to deliver a functional load of lipids, emollients, sunscreens, liquid silicone oils and even water. Particularly attractive would be a capability to incorporate an aqueous emulsion into the powder system. Water soluble compounds could then more readily be incorporated into powders normally unfriendly to such compounds.

35

Accordingly, it is an object of the present invention to provide a cosmetic powder which delivers improved

- 2 -

aesthetics while greatly improving functionality and skinfeel.

5 Another object of the present invention is to provide a cosmetic powder wherein water can be incorporated and thereby also incorporate water soluble compounds.

10 These and other objects of the present invention will become more apparent from consideration of the following summary and description.

#### SUMMARY OF THE INVENTION

15 A cosmetic powder is provided including:

- (i) from 0.1 to 50% of a crosslinked non-emulsifying siloxane elastomer;
- 20 (ii) from 1 to 99% of a powdered inorganic material; and
- (iii) from 0.1 to 20% of a skin treatment agent.

25 Inorganic powders can now be formulated with high levels of hydrophobic and/or hydrophilic liquids through coupling action of the crosslinked non-emulsifying siloxane elastomer. For instance, water can be incorporated into a hydrophobic liquid coupled with the elastomer which is then loaded onto the inorganic powder. Water soluble actives

30 such as the salts of alpha-hydroxycarboxylic acids can be dissolved in the incorporated water. Thereby normally incompatible water soluble salts can be folded into a hydrophobic system which itself is locked within a cosmetic powder.

35

- 3 -

DETAILED DESCRIPTION OF THE INVENTION

Now it has been discovered that crosslinked non-emulsifying siloxane elastomers can serve as a coupling vehicle for introducing skin treatment agents into a powder formed mainly of inorganic material.

Crosslinked non-emulsifying siloxane elastomers are a first essential element of this invention. They will usually have an average number molecular weight in excess of 2,000, preferably in excess of 1,000,000. Typically they will range from 10,000 to 20 million. The term "non-emulsifying" defines a siloxane from which polyoxyalkylene units are absent. Advantageously the elastomers are formed from a divinyl compound, particularly a siloxane polymer with at least two free vinyl groups, reacting with Si-H linkages of a polysiloxane backbone such as a molecularly spherical MQ resin. Suitable elastomer compositions are commercially available from the General Electric Company under product designation General Electric Silicone 1229 with proposed CTFA name of Cyclomethicone and Vinyl Dimethicone/Methicone Cross Polymer, delivered as 20-35% elastomer in a cyclomethicone carrier. A related elastomer composition under the CTFA name of Crosslinked Stearyl Methyl Dimethyl Siloxane Copolymer is available as Gransil SR-CYC (25-35% elastomer in cyclomethicone) from Grant Industries, Inc., Elmwood Park, New Jersey. Elastomer without cyclomethicone is available from Dow Corning as DC 9506. The commercial products from General Electric and Grant Industries preferably are further processed by subjecting them to a high pressure (approximately 5,000 psi) treatment in a Sonolator with recycling in 10 to 60 passes. Sonolation achieves a resultant fluid with elastomer average particle size ranging from 0.2 to 10 micron, preferably 0.5 to 5 micron. Viscosity is best when

- 4 -

ranging between 300 and 20,000 cps at 25°C as measured by a Brookfield LV Viscometer (size 4 bar, 60 rpm, 15 sec.).

5 Amounts of the elastomer may range from 0.1 to 50%,  
optimally from 1 to 15%, most preferably from 3 to 10% by  
weight of the composition.

10 A second essential element of the present invention is that  
of a powdered inorganic material. The powdered material is  
preferably chosen from talc, mica, mineral silicates and  
mixtures thereof.

15 Micas useful in the present invention are in particular  
muscovite, phlogopite, tiotite, sericite, lepidolite,  
paragonite and artificial or synthetic mica having a  
fluorine atom substituted for the hydroxyl group of natural  
mica as well as baked or calcined products thereof. Such  
micas may be used alone or in any mixture thereof.

20 Mineral silicates useful in the present invention are for  
example phyllosilicates and tectosilicates such as  
pyrophyllite, chlorite, antigorite, lizardite, kaolinite,  
dickite, nacrite, halloysite, montmorillonite, nontronite,  
saponite, sauconite, and bentonite; natrolites such as  
25 natrolite, mesolite, scolecite, and thomsonite; heulandites  
such as heulandite, stilbite, epistibite; and zeolites such  
as analcite, harmotome, phillipsite, chabazite and  
gmelinite. Such silicate minerals may be used alone or in  
combination. The phyllosilicates may have organic cations  
30 at the interface of the layers thereof or may be  
substituted with alkali metal or alkaline earth metal ions.  
The tectosilicates may include metallic ions in the fine  
pores thereof.

- 5 -

Amounts of the powdered inorganic material may range from 1 to 99%, preferably from 30 to 85%, optimally from 50 to 75% by weight.

5 Average particle size of powdered inorganic material according to the present invention preferably ranges from 1 to 1,000 microns more preferably from 5 to 600 microns, optimally from 80 to 300 microns. The fully formulated cosmetic powder compositions of the present invention  
10 preferably has will have similar average particle sizes.

A third element of the present invention is that of a skin treatment agent. This agent is preferably selected from the group consisting of moisturizing conditioners,  
15 exfoliants, sunscreens, deodorants and mixtures thereof.

Moisturizing conditioners include humectants, hydrocarbons and oily ester emollients. Humectants are usually polyhydric alcohols. Not only do they moisturize/condition  
20 but they can function to reduce scaling and stimulate removal of built-up scale from the skin. Typical polyhydric alcohols include polyalkylene glycols and more preferably alkylene polyols and their derivatives. Illustrative are propylene glycol, dipropylene glycol,  
25 polypropylene glycol, polyethylene glycol, sorbitol, hydroxypropyl sorbitol, hexylene glycol, 1,3-butylene glycol, 1,2,6-hexanetriol, ethoxylated glycerin, propoxylated glycerin and mixtures thereof. Most preferably the humectant is glycerin. The moisturizing  
30 conditioners may also include emollients which can be selected from hydrocarbons or esters. Petrolatum is the most preferred conditioner. Other hydrocarbons that may be employed include mineral oil, polyolefins such as polydecene, and parafins such as isohexadecane (e.g.  
35 Permethyl 99® and Permethyl 101®).

- 6 -

Oily ester emollients which moisturize may be those selected from one or more of the following classes:

- 5 1. Triglyceride esters such as vegetable and animal fats and oils. Examples include castor oil, cocoa butter, safflower oil, cottonseed oil, corn oil, olive oil, cod liver oil, almond oil, avocado oil, palm oil, sesame oil, squalene, Kikui oil and soybean oil.
- 10 2. Acetoglyceride esters, such as acetylated monoglycerides.
- 15 3. Ethoxylated glycerides, such as ethoxylated glyceryl monostearate.
- 20 4. Alkyl esters of fatty acids having 10 to 20 carbon atoms. Methyl, isopropyl, and butyl esters of fatty acids are useful herein. Examples include hexyl laurate, isohexyl laurate, isohexyl palmitate, isopropyl palmitate, decyl oleate, isodecyl oleate, hexadecyl stearate, decyl stearate, isopropyl isostearate, diisopropyl adipate, diisohexyl adipate, dihexyldecyl adipate, diisopropyl sebacate, lauryl lactate, myristyl lactate, and cetyl lactate.
- 25 5. Alkenyl esters of fatty acids having 10 to 20 carbon atoms. Examples thereof include oleyl myristate, oleyl stearate, and oleyl oleate.
- 30 6. Ether-esters such as fatty acid esters of ethoxylated fatty alcohols.
- 35 7. Polyhydric alcohol esters. Ethylene glycol mono and di-fatty acid esters, diethylene glycol mono- and di-fatty acid esters, polyethylene glycol (200-6000) mono- and di-fatty acid esters, propylene glycol mono-



- 7 -

and di-fatty acid esters, polypropylene glycol 2000 monooleate, polypropylene glycol 2000 monostearate, ethoxylated propylene glycol monostearate, glyceryl mono- and di-fatty acid esters, polyglycerol polyfatty esters, ethoxylated glyceryl monostearate, 1,2-butylene glycol monostearate, 1,2-butylene glycol distearate, polyoxyethylene polyol fatty acid ester, sorbitan fatty acid esters, and polyoxyethylene sorbitan fatty acid esters are satisfactory polyhydric alcohol esters.

8. Wax esters such as beeswax, spermaceti, myristyl myristate, stearyl stearate.

Exfoliants according to the present invention are preferably selected from alpha-hydroxycarboxylic acids, beta-hydroxycarboxylic acids and salts of these acids. Most preferred are glycolic, lactic and salicylic acids and their ammonium salts.

A wide variety of alpha-hydroxycarboxylic acids, preferably C<sub>2</sub>- C<sub>30</sub> alpha-hydroxycarboxylic acids, may be employed. Suitable examples of which include:

- hydroxyethanoic acid
- hydroxypropanoic acid
- hydroxyhexanoic acid
- hydroxyoctanoic acid
- hydroxydecanoic acid
- hydroxydodecanoic acid
- hydroxytetradecanoic acid
- hydroxyhexadecanoic acid
- hydroxyoctadecanoic acid
- hydroxyeicosanoic acid
- hydroxydocosanoic acid
- hydroxyhexacosanoic acid, and
- hydroxyoctacosanoic acid

- 8 -

Sunscreens may also be used as the skin treatment agent. The sunscreens are preferably selected from 2-ethylhexyl p-methoxycinnamate, 4,4'-t-butyl methoxydibenzoylmethane, 2-hydroxy-4-methoxybenzophenone, octyldimethyl p-aminobenzoic acid, digalloyltriolate, 2,2-dihydroxy-4-methoxybenzophenone, ethyl-4-[bis(hydroxypropyl)]aminobenzoate, 2-ethylhexyl-2-cyano-3,3-diphenylacrylate, 2-ethylhexylsalicylate, glyceryl p-aminobenzoate, 3,3,5-trimethylcyclohexylsalicylate, methylanthranilate, p-dimethylaminobenzoic acid or aminobenzoate, 2-ethylhexyl p-dimethylaminobenzoate, 2-phenylbenzimidazole-5-sulfonic acid, 2-(p-dimethylaminophenyl)-5-sulfoniobenzoxazoic acid and mixtures thereof.

Deodorants should preferably be capable of killing or hindering the growth of microorganisms that generate malodour or that promote the decomposition of body oils into odiferous fatty acids. Most prominent among organic antimicrobial materials are triclosan, triclorban, chlorhexedine, dibromodicyanobutane, 2-bromo-2-nitropropane-diol-1,3, octenidine salts, alexidine salts, and certain fragrant oils known as deo perfumes (e.g. U.S. Patent 4,278,658 to Hooper et al.). Inorganic antimicrobial materials may also serve as deodorant actives. These include zinc oxide, zinc hydroxide, zinc carbonate, zinc phenolsulfonate, magnesium oxide, magnesium hydroxide, magnesium carbonate, lanthanum oxide, lanthanum hydroxide, lanthanum carbonate and combinations thereof.

Deodorancy may also be achieved from astringent salts such as those of aluminum, zirconium, zinc and mixtures thereof. Salts useful as astringents or as components of astringent complexes include aluminum halides, aluminum hydroxyhalides, zirconyl oxyhalides, zirconyl hydroxyhalides and mixtures of these salt materials.

- 9 -

Amounts of the skin treatment agent may range from 0.1 to 20%, preferably from 0.5 to 15%, optimally from 1 to 10% by weight. Particularly preferred skin treatment agents are  $\alpha$ -hydroxycarboxylic acids,  $\beta$ -hydroxycarboxylic acids and salts thereof.

Compositions of this invention may include volatile and nonvolatile silicones other than the crosslinked elastomers.

The term "volatile" refers to those materials having a measurable pressure at ambient conditions. Volatile polyorganosiloxanes useful herein may be cyclic or linear. Preferred cyclic silicones include polydimethylsiloxanes containing from about 3 to about 9 silicon atoms, preferably containing from about 4 to about 5 silicon atoms, generally known as cyclomethicones. Preferred linear silicone oils include the polydimethylsiloxanes containing from about 3 to about 9 silicon atoms. The linear volatile silicones generally have viscosities of less than about 5 centistokes at 25°C, while the cyclic materials have viscosities of less than about 10 centistokes, the preferable range being from 0.1 to 8 centistokes. Examples of silicone oils useful in the present invention include: Dow Corning 344, Dow Corning 345 and Dow Corning 244 (manufactured by the Dow Corning Corporation); Silicone 7207 and Silicone 7158 (manufactured by the Union Carbide Corporation); SF1202 (manufactured by General Electric).

Non-volatile polyorganosiloxanes may be fluids selected from polyalkyl siloxane, polyalkylaryl siloxane or polyether siloxane copolymers. The non-volatile polyalkyl siloxanes that may be used include, for example, polydimethyl siloxanes with viscosities ranging from about 5 to 100,000 centistokes at 25°C. These siloxanes are

- 10 -

available from the General Electric Company as the Vicasil series and from Dow Corning as the Dow Corning 200 series. The non-volatile polyalkylaryl siloxanes may include, for example, polymethylphenylsiloxanes having viscosities of from 15 to 65 centistokes at 25°C. These siloxanes are available from the General Electric Company as SF 1075 methyl phenyl fluid or from Dow Corning as 556 Cosmetic Grade Fluid. The non-volatile polyether siloxane copolymer may include, for example, a dimethyl polyoxyalkylene ether copolymer fluid having a nominal viscosity of about 1200 to 1500 centistokes at 25°C. The copolymer is available from the General Electric Company as SF-1066 organosilicone surfactant. Preferred compounds of this type are polysiloxane ethylene glycol ether copolymers.

Cosmetic powders of the present invention are preferably prepared in the following manner. An aqueous emulsion is formed from a combination of water and a volatile oil siloxane elastomer blend. One or more skin treatment agents are emulsified or otherwise incorporated into the emulsion. It is most preferable that a water-in-oil type emulsion be formed. When the aqueous phase is internal, the phase will later be protected from evaporation and thereby entrapped with the dried residue of the crosslinked elastomer. Volatile oils such as cyclomethicone and C<sub>5</sub>-C<sub>10</sub> hydrocarbons are useful as the evaporative component. Relative weight ratios of the water and oil phases may range from 20:1 to 1:2, preferably from 10:1 to 1:1, optimally from 6:1 to 2:1. Subsequent to emulsion formation, volatile oils are stripped by evaporation. The resultant dried blend is then mixed with powdered inorganic material such as talc. The resultant powder is then subjected to high shear mixing in an Osterizer or equivalent apparatus.

- 11 -

The following examples will more fully illustrate the embodiments of this invention. All parts, percentages and proportions referred to herein and in the appended claims are by weight unless otherwise indicated.

**EXAMPLE 1**

This Example illustrates a typical talcum powder product achievable by the present invention. A water-in-oil emulsion is formed by blending the potassium lactate solution into elastomer dispersed within cyclomethicone (with petrolatum and emulsifier). The resultant blend (see Table I) is then poured into and mixed with powdered talc in a ratio of 1:10 to form a talcum powder.

**TABLE I**

COMPONENT	WEIGHT %
Gransil SR-CYC7	22.5
DC 3447 (Cyclomethicone)	54
Petrolatum	11
Potassium Lactate (50% Water)	7
ABIL EM907 (Silicone copolyol)	0.5

**EXAMPLE 2**

This Example provides a series of comparative experiments to demonstrate the unique character of the silicone elastomer in contrast to other thickening silicone materials. Table II lists the formulations and Table III describes the physical properties of the resultant compositions.

TABLE II

TRADE NAME	CTFA NAME	FORMULATIONS (WEIGHT %)					
		A	B	C	D	E	F
PJ 2.5 hard	Petrolatum	0.3	0.3	0.3	0.3	0.3	0.3
Purac BFP39	Potassium Lactate	5	5	5	5	5	5
Permethyl 99a	C12-14 Isoparaffin	0.5	0.5	0.5	0.5	0.5	0.5
Abil EM90	Cetyl dimethicone-copolyol	0.2	0.2	0.2	0.2	0.2	0.2
Talc	Talc	54	54	54	54	54	54
DC 9506	Silicone Elastomer	40					
DC 593	Trimethylsiloxysilicate		40				
Abil 9801	Cetyl-Dimethicone			40			
Abil 9800	Stearyl Dimethicone				40		
GE SE30	Dimethicone					40	
Dow 556	Phenyl Trimethicone						40

- 13 -

**TABLE III**

	FORMULATIONS					
	A	B	C	D	E	F
Results	Flowable powder	Very thick pourable	Pourable Liquid	Solid non-powder	Slightly pourable very thick liquid	Pourable liquid

Based upon the results in Table III, only the silicone elastomer (Dow Corning 9506) provided a composition that was a flowable powder. All other formulations were not in powder form.

**CLAIMS**

1. A cosmetic powder comprising:
  - (i) from 0.1 to 50% of a crosslinked non-emulsifying siloxane elastomer;
  - (ii) from 1 to 99% of a powdered inorganic material; and
  - (iii) from 0.1 to 20% of a skin treatment agent.
2. A powder according to claim 1 wherein the skin treatment agent is selected from the group consisting of moisturizing conditioners, exfoliants, sunscreens, deodorants and mixtures thereof.
3. A powder according to claim 1 or claim 2 wherein the skin treatment agent is selected from the group consisting of alpha-hydroxycarboxylic acid, beta-hydroxycarboxylic acid and salts thereof.
4. A powder according to any one of claims 1-3 wherein the crosslinked non-emulsifying siloxane elastomer is formed from a divinyl compound reacted with Si-H linkages of a polysiloxane.
5. A powder according to any one of claims 1-4 wherein the inorganic material is talc.



# INTERNATIONAL SEARCH REPORT

International Application No.  
PCT/EP 97/03063

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 6 A61K7/48 A61K7/035

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol. 16, no. 156 (C-930) & JP 04 009323 A (KOBAYASHI KOSE) see abstract	1-5
A	STN, File Supplier, Karlsruhe, DE, File XP002039716 Chemical Abstracts, vol 115, AN=286965 see the abstract & JP 03 197 413 A	1-5
A	STN, File Supplier, Karlsruhe, DE, File XP002039717 Chemical Abstracts, vol 115, AN=15313 see the abstract & JP 02 304 015 A	1-5

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

### \* Special categories of cited documents:

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

- \*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- \*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- \*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- \*A\* document member of the same patent family

Date of the actual completion of the international search

4 September 1997

Date of mailing of the international search report

19. 09. 97

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax (+31-70) 340-3016

Authorized officer

Fischer, J.P.

# INTERNATIONAL SEARCH REPORT

Int'l Application No  
PCT/EP 97/03063

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	FR 2 688 134 A (L'OREAL) 10 September 1993 see the whole document ---	1-5
A	EP 0 709 083 A (REVLON CONSUMER PRODUCTS) 1 May 1996 see the whole document ---	1-5
A	WO 95 25499 A (PROCTER & GAMBLE) 28 September 1995 see the whole document ---	1-5
A	EP 0 388 582 A (KOBAYASHI KOSE) 26 September 1990 see the whole document -----	1-5

# INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/EP 97/03063

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
FR 2688134 A	10-09-93	CA 2102189 A	06-09-93
		DE 69304162 D	26-09-96
		DE 69304162 T	13-03-97
		EP 0583460 A	23-02-94
		ES 2093412 T	16-12-96
		WO 9317660 A	16-09-93
		JP 6507426 T	25-08-94
		US 5496544 A	05-03-96
-----			
EP 709083 A	01-05-96	AU 3439195 A	09-05-96
		BR 9504533 A	27-05-97
		CA 2161285 A	26-04-96
		CN 1127631 A	31-07-96
		FI 955048 A	26-04-96
		FR 2726467 A	10-05-96
		GB 2294392 A	01-05-96
		JP 8239316 A	17-09-96
		NO 954201 A	26-04-96
		NZ 280302 A	20-12-96
		ZA 9509024 A	06-08-96
		-----	
WO 9525499 A	28-09-95	AU 1934195 A	09-10-95
		CA 2184907 A	28-09-95
		CN 1143905 A	26-02-97
		EP 0785763 A	30-07-97
-----			
EP 388582 A	26-09-90	JP 2247110 A	02-10-90
		JP 2250812 A	08-10-90
		JP 2132141 A	21-05-90
		JP 2258709 A	19-10-90
		JP 2279617 A	15-11-90
		DE 69014358 D	12-01-95
		DE 69014358 T	14-06-95
		US 5061481 A	29-10-91
		US 5219560 A	15-06-93
		JP 3008432 A	16-01-91
-----			